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- 1. A substantially purified integral membrane protein comprising the amino acid sequence of SEQ ID NO:1 or fragments thereof.
- 2. An isolated and purified polynucleotide sequence encoding the integral membrane protein of claim 1.
- 3. A polynucleotide sequence which hybridizes under stringent conditions to the polynucleotide sequence of claim 2.
 - 4. A composition comprising the polynucleotide sequence of claim 2.
- 5. An isolated and purified polynucleotide sequence comprising SEQ ID NO:2 or variants thereof.
- 6. A polynucleotide sequence which is complementary to SEQ ID NO:2 or variants thereof.
 - 7. A composition comprising the polynucleotide sequence of claim 6.
 - 8. An expression vector containing the polynucleotide sequence of claim 2.
 - 9. A host cell containing the vector of claim 8.
- 10. A method for producing a polypeptide comprising the amino acid sequence of SEQ ID NO:1 or fragments thereof, the method comprising the steps of:
 - a) culturing the host cell of claim 9 under conditions suitable for the

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expression of the polypeptide; and

- recovering the polypeptide from the host cell culture. b)
- A pharmaceutical composition comprising a substantially purified human 11. IMP-2 protein having an amino acid sequence of SEQ ID NO:1/in conjunction with a suitable pharmaceutical carrier.
 - A purified antibody which binds specifically to the polypeptide of claim 1. 12.
 - A purified agonist which specifically binds to and modulates the activity of the 13. polypeptide of claim 1.
- A purified antagonist which specifically binds to and modulates the activity of 14. the polypeptide of claim 1.
 - A pharmaceutical composition comprising the antagonist of claim 14. 15.
- 16. A method for treating liver disease comprising administering to a subject in need of such treatment an effective amount of the pharmaceutical composition of claim 13.
- A method for detection of polynucleotides encoding human IMP-2 in a biological sample comprising the steps of:
 - hybridizing the polynucleotide of claim 2 to nucleic acid material of a biological sample, the
 - detecting said hybridization complex, wherein the presence of said complex correlates with the presence of a polynucleotide encoding human IMP-2 in said biological sample.
 - 18. The method of claim 17, wherein before hybridization, the nucleic acid

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material of the biological sample is amplified by the polymerase chain reaction.